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(FILE 'HOME' ENTERED AT 21:16:46 ON 10 MAR 2009)
     FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH, LIFESCI' ENTERED AT 21:17:27 ON
     10 MAR 2009
            417 S (OSTEOBLAST? OR OSTEOGEN? OR STEM(W)CELL)(7A)DEDIFFERENTIAT?
L2
         286805 S ALKALINE (W) PHOSPHATASE OR COLLAGEN (W) TYPE (3A) I OR OSTEOCALCIN
L3
           2958 S (SUBSTITUT? OR DISUBSTITUT?) (4A) PURINE
L4
             24 S L1(P)L2
L5
             0 S L3 AND L4
L6
             0 S L1 AND L2 AND L3
             1 S L1 AND L3
L8
             10 DUP REM L4 (14 DUPLICATES REMOVED)
L9
            523 S (ADIPOCYTE OR ADIPOGEN? OR STEM(W) CELL) (7A) DEDIFFERENTIAT?
L10
          79926 S OB OR UCP OR PPARGAMMA OR C/EBP
L11
         98923 S OB OR UCP OR PPARGAMMA OR C(W) EBP
L12
             33 S L9(P)L11
             10 DUP REM L12 (23 DUPLICATES REMOVED)
L13
L14
              0 S L3 AND L12
=> d bib ab 17
     ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN
ΑN
     2003:996204 CAPLUS
DN
     140:160988
ΤТ
     Dedifferentiation of Lineage-Committed Cells by a Small Molecule
AU
     Chen, Shuibing; Zhang, Qisheng; Wu, Xu; Schultz, Peter G.; Ding, Sheng
CS
     Department of Chemistry and the Skaggs Institute for Chemical Biology, The
     Scripps Research Institute, La Jolla, CA, 92037, USA
SO
     Journal of the American Chemical Society (2004), 126(2), 410-411
     CODEN: JACSAT; ISSN: 0002-7863
PB
    American Chemical Society
     Journal
DT
LA
    English
AB
    Combinatorial libraries were screened for mols. that induce mouse myogenic
     lineage committed cells to dedifferentiate in vitro. A 2,6-
     disubstituted purine, reversine, was discovered that
     induces lineage reversal of C2C12 cells to become multipotent progenitor
     cells which can redifferentiate into osteoblasts and adipocytes. This and
     other such mols, are likely to provide new insights into the mol.
     mechanisms that control cellular dedifferentiation and may ultimately be
     useful to in vivo stem cell biol, and therapy,
RE.CNT 12
             THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
=> d au ti so pi 1-10 18
     ANSWER 1 OF 10 CAPLUS COPYRIGHT 2009 ACS on STN
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L8

Kudo, Naoko; Ogose, Akira; Hotta, Tetsuo; Kawashima, Hiroyuki; Gu, Wenguang; Umezu, Hajime; Toyama, Tsuyoshi; Endo, Naoto ΑU

Establishment of novel human dedifferentiated chondrosarcoma cell line with osteoblastic differentiation

Virchows Archiv (2007), 451(3), 691-699 CODEN: VARCEM; ISSN: 0945-6317

1.8 ANSWER 2 OF 10 LIFESCI COPYRIGHT 2009 CSA on STN

AU Steck, Eric; Bertram, Helge; Abel, Rainer; Chen, Bohua; Winter, Anja; Ritcher, Wiltrud

Induction of Intervertebral Disc-Like Cells From Adult Mesenchymal Stem

Cells

SO. Stem Cells, (20050300) vol. 23, no. 3, pp. 403-411. ISSN: 1066-5099.

- ANSWER 3 OF 10 MEDLINE on STN T.R DUPLICATE 1
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- Exogenous Runx2 expression enhances in vitro osteoblastic differentiation TT
- and mineralization in primary bone marrow stromal cells.
- SO Tissue engineering, (2004 Nov-Dec) Vol. 10, No. 11-12, pp. 1623-32. Journal code: 9505538, ISSN: 1076-3279.
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- TТ Vascular calcification in chronic kidney disease.
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- ΤI
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DUPLICATE 2

DUPLICATE 3

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 Journal code: 7802877. ISSN: 0021-9738.
 Report No.: NLM-PMC442840.

=> d ab 4 113

L13 ANSWER 4 OF 10 MEDLINE on STN DUPLICATE 2

obesity, including type II diabetes.

AB This study identifies monocyte chemoattractant protein 1 (MCP-1) as an insulin-responsive gene. It also shows that insulin induces substantial expression and secretion of MCP-1 both in vitro in insulin-resistant (IR) 3T3-L1 adipocytes and in vivo in IR obese mice (ob/ob). Thus, MCP-1 resembles other previously described genes (e.g., PAI-1 and SREBP-1c) that remain sensitive to insulin in IR states. The hyperinsulinemia that frequently accompanies obesity and insulin resistance may therefore contribute to the altered expression of these and other genes in insulin target tissues. In vivo studies also demonstrate that MCP-1 is overexpressed in obese mice compared with their lean controls, and that white adipose tissue is a major source of MCP-1. The elevated MCP-1 may alter adipocyte function because addition of MCP-1 to differentiated adipocytes in vitro decreases insulin-stimulated glucose uptake and the expression of several adipogenic genes (LpL, adipsin, GLUT-4, aP2, beta3-adrenergic receptor, and peroxisome proliferator-activated receptor gamma). These results suggest that elevated MCP-1 may induce adipocyte dedifferentiation and contribute to pathologies associated with hyperinsulinemia and